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NEWS 3 SEP 09 ACD predicted properties enhanced in REGISTRY/ZREGISTRY
NEWS 4 OCT 03 MATHDI removed from STN
NEWS 5 OCT 04 CA/CAPLUS-Canadian Intellectual Property Office (CIPO) added
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NEWS 6 OCT 13 New CAS Information Use Policies Effective October 17, 2005
NEWS 7 OCT 17 STN(R) AnaVist(TM), Version 1.01, allows the export/download
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NEWS 10 OCT 27 EPFULL enhanced with additional content
NEWS 11 NOV 14 CA/CAPLUS - Expanded coverage of German academic research
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FILE 'HOME' ENTERED AT 11:27:39 ON 07 DEC 2005

=> index bioscience

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COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
0.21	0.21

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INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, ANTE, AQUALINE,
AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS,
BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB,
CROPU, DDFB, DDFU, DGENE, DISSABS, ...' ENTERED AT 11:27:52 ON 07 DEC 2005

74 FILES IN THE FILE LIST IN STNINDEX

Enter SET DETAIL ON to see search term postings or to view
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=> (fluorescent or chromoprotein) and emission and maximum

23 FILE AGRICOLA

96 FILE ANABSTR
 1 FILE ANTE
 9 FILE AQUALINE
 53 FILE AQUASCI
 18 FILE BIOBUSINESS
 46 FILE BIOENG
 824 FILE BIOSIS
 70 FILE BIOTECHABS
 70 FILE BIOTECHDS
 207 FILE BIOTECHNO
 64 FILE CABA
 43 FILE CANCERLIT
 2381 FILE CAPLUS
 7 FILE CEABA-VTB
 7 FILE CEN
 1 FILE CIN
 1 FILE CONFSCI
 8 FILE CROPU
 9 FILE DDFU
 594 FILE DGENE
 84 FILE DISSABS

29 FILES SEARCHED...

72 FILE DRUGU
 5 FILE EMBAL
 589 FILE EMBASE
 254 FILE ESBIODBASE
 9 FILE FEDRIP
 7 FILE FROSTI
 16 FILE FSTA
 2 FILE HEALSAFE
 618 FILE IFIPAT
 135 FILE JICST-EPLUS
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 774 FILE MEDLINE
 22 FILE NIOSHTIC
 27 FILE NTIS
 19 FILE OCEAN
 361 FILE PASCAL

57 FILES SEARCHED...

3 FILE PHIN
 197 FILE PROMT
 22 FILE RDISCLOSURE
 878 FILE SCISEARCH
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 22952 FILE USPATFULL
 2659 FILE USPAT2
 2 FILE VETU
 21 FILE WATER
 452 FILE WPIDS
 1 FILE WPIFV
 452 FILE WPINDEX

51 FILES HAVE ONE OR MORE ANSWERS, 74 FILES SEARCHED IN STNINDEX

L1 QUE (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM

=> d rank

F1 22952 USPATFULL
 F2 2659 USPAT2
 F3 2381 CAPLUS
 F4 878 SCISEARCH
 F5 824 BIOSIS
 F6 774 MEDLINE

F7	618	IFIPAT
F8	594	DGENE
F9	589	EMBASE
F10	452	WPIDS
F11	452	WPINDEX
F12	361	PASCAL
F13	254	ESBIOBASE
F14	228	TOXCENTER
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F16	197	PROMT
F17	168	LIFESCI
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F19	96	ANABSTR
F20	84	DISSABS
F21	72	DRUGU
F22	70	BIOTECHABS
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F24	64	CABA
F25	53	AQUASCI
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F27	43	CANCERLIT
F28	27	NTIS
F29	23	AGRICOLA
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F32	21	WATER
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F35	16	FSTA
F36	9	AQUALINE
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F38	9	FEDRIP
F39	8	CROPU
F40	7	CEABA-VTB
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F42	7	FROSTI
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F44	3	PHIN
F45	2	HEALSAFE
F46	2	VETU
F47	1	ANTE
F48	1	CIN
F49	1	CONFSCI
F50	1	KOSMET
F51	1	WPIFV

=> file biosis medline caplus
COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
2.95	3.16

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=> (fluorescent or chromoprotein) and emission and maximum and nm
L2 2641 (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM AND NM

=> (fluorescent or chromoprotein) and emission and maximum and "far red shifted"
L3 1 (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM AND
"FAR RED SHIFTED"

=> d ab bib

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN
AB Nucleic acid compns. encoding Stichodactylidaen **chromoproteins**
and **fluorescent** mutants thereof, as well as the polypeptide
compns. encoded by the same, are provided. The proteins of interest are
proteins that are colored and/or **fluorescent**, where this feature
arises from the interaction of two or more residues of the protein. Also
of interest are proteins that are substantially similar to, or mutants of,
the above specific proteins, including non-aggregating mutants and mutants
with modulated oligomerization characteristics as compared to wild type.
Thus, the two wild-type **chromoprotein** isoforms from Heteractis
crispa exhibit a strong **emission max.** at
.apprx.580-640 nm. Site-specific mutagenesis of the Cys-148 residue to
serine dramatically increases the quantum yield of red fluorescence as
compared to the wild-type protein, and further random mutagenesis (A2S,
T36A, C143S, L173H, P201L, K204E) yielded an even brighter mutant. A
single mutation, L126H, may be responsible for modifying the oligomeric
state of the protein from tetrameric to dimeric. Also provided are
fragments of the nucleic acids and the peptides encoded thereby, as well
as antibodies to the subject proteins and transgenic cells and organisms.
The subject protein and nucleic acid compns. find use in a variety of
different applications. Finally, kits for use in such applications, e.g.,
that include the subject nucleic acid compns., are provided.

AN 2002:293685 CAPLUS

DN 136:321045

TI **Far red-shifted fluorescent**

proteins and their variants from Heteractis crispis

IN Lukyanov, Sergey Anatolievich; Fradkov, Arcady Fedorovich; Lukyanov,
Konstantin Anatolievich; Gurskaya, Nadezda Georgievna

PA Clontech Laboratories, Inc., USA

SO PCT Int. Appl., 81 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 17

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-----	---	-----	-----	-----
PI	WO 2002030965	A2	20020418	WO 2001-US32080	20011012
	WO 2002030965	A3	20030605		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,				
	CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,				
	GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,				
	LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL,				
	PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,				
	US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG,				
	KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR,				
	IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN,				
	GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2425318	AA	20020418	CA 2001-2425318	20011012
	AU 2002011722	A5	20020422	AU 2002-11722	20011012
	EP 1334122	A2	20030813	EP 2001-979797	20011012
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,				
	IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	JP 2004532606	T2	20041028	JP 2002-534350	20011012
PRAI	US 2000-240018P	P	20001012		
	US 2001-306131P	P	20010716		
	WO 2001-US32080	W	20011012		

=> (fluorescent or chromoprotein) and emission and maximum and " red shifted"
L4 124 (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM AND "
RED SHIFTED"

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ENTER L# LIST OR (END):14
PROCESSING COMPLETED FOR L4
L5 82 DUP REMOVE L4 (42 DUPLICATES REMOVED)

=> 15 and nm
L6 56 L5 AND NM

=> dup remove
ENTER L# LIST OR (END):16
PROCESSING COMPLETED FOR L6
L7 56 DUP REMOVE L6 (0 DUPLICATES REMOVED)

=> 16 and nucleic with acid
L8 1 L6 AND NUCLEIC WITH ACID

=> d ab bib

L8 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN
AB **Nucleic acid** compns. encoding Stichodactylidaen
chromoproteins and **fluorescent** mutants thereof, as well
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DN 136:321045

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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

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PRAI US 2000-240018P	P	20001012		
US 2001-306131P	P	20010716		
WO 2001-US32080	W	20011012		

=> 16 and" nucleic acid"
L9 1 L6 AND" NUCLEIC ACID"

=> d ab bib

L9 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN

AB **Nucleic acid** compns. encoding Stichodactylidaen **chromoproteins** and **fluorescent** mutants thereof, as well as the polypeptide compns. encoded by the same, are provided. The proteins of interest are proteins that are colored and/or **fluorescent**, where this feature arises from the interaction of two or more residues of the protein. Also of interest are proteins that are substantially similar to, or mutants of, the above specific proteins, including non-aggregating mutants and mutants with modulated oligomerization characteristics as compared to wild type. Thus, the two wild-type **chromoprotein** isoforms from Heteractis crispa exhibit a strong **emission max.** at .apprx.580-640 nm. Site-specific mutagenesis of the Cys-148 residue to serine dramatically increases the quantum yield of red fluorescence as compared to the wild-type protein, and further random mutagenesis (A2S, T36A, C143S, L173H, P201L, K204E) yielded an even brighter mutant. A single mutation, L126H, may be responsible for modifying the oligomeric state of the protein from tetrameric to dimeric. Also provided are fragments of the **nucleic acids** and the peptides encoded thereby, as well as antibodies to the subject proteins and transgenic cells and organisms. The subject protein and **nucleic acid** compns. find use in a variety of different applications. Finally, kits for use in such applications, e.g., that include the subject **nucleic acid** compns., are provided.

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TI Far **red-shifted fluorescent** proteins and their variants from Heteractis crispis

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	WO 2002030965	A3	20030605		
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	RW:		GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG		
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PRAI	US 2000-240018P	P	20001012		
	US 2001-306131P	P	20010716		
	WO 2001-US32080	W	20011012		

=> l6 and anthozoan

L10 0 L6 AND ANTHOZOAN

=> l6 and polypeptide

L11 1 L6 AND POLYPEPTIDE

=> l5 and anthozoan

L12 0 L5 AND ANTHOZOAN

=> (fluorescent or chromoprotein) and emission and maximum and " red shifted" and anthozoan

L13 0 (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM AND " RED SHIFTED" AND ANTHOZOAN

=> d l6 ti 1-20

L6 ANSWER 1 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Two-photon excitation and **emission** spectra of the green **fluorescent** protein variants ECFP, EGFP and EYFP.

L6 ANSWER 2 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI The two photon cross section and spectrum of 6MAP, a **fluorescent** adenosine analog.

L6 ANSWER 3 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Crystallization and preliminary X-ray diffraction analysis of the red **fluorescent** protein eqFP611.

L6 ANSWER 4 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Steady-state and time-resolved fluorescence studies indicate an unusual conformation of 2-aminopurine within ATAT and TATA duplex DNA sequences.

L6 ANSWER 5 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI A far-red **fluorescent** protein with fast maturation and reduced oligomerization tendency from Entacmaea quadricolor (Anthozoa, Actinaria).

L6 ANSWER 6 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Kinetic analysis of maturation and denaturation of DsRed, a coral-derived red **fluorescent** protein.

L6 ANSWER 7 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN

TI Suitability of enhanced green fluorescent protein as a reporter component for bioassays.

L6 ANSWER 8 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI The structure of the chromophore within DsRed, a red fluorescent protein from coral.

L6 ANSWER 9 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI Novel fluorescent protein from *Discosoma coral* and its mutants possesses a unique far-red fluorescence.

L6 ANSWER 10 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI Fluorescent probing of membrane potential in walled cells: diS-C3(3) assay in *Saccharomyces cerevisiae*.

L6 ANSWER 11 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI Terminal Marking of Triosephosphate Isomerase: Consequences of Deamidation.

L6 ANSWER 12 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI Quenching of intrinsic fluorescence of yeast cytochrome c peroxidase by covalently- and noncovalently-bound quenchers.

L6 ANSWER 13 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI Low temperature absorbance and fluorescence spectroscopy of the photoactive yellow protein from *Ectothiorhodospira halophila*.

L6 ANSWER 14 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI PICOSECOND DECAY KINETICS AND QUANTUM YIELD OF FLUORESCENCE OF THE PHOTOACTIVE YELLOW PROTEIN FROM THE HALOPHILIC PURPLE PHOTOTROPHIC BACTERIUM *ECTOTHIORHODOSPIRA-HALOPHILA*.

L6 ANSWER 15 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI INTERACTION OF MUSCLE AND NON-MUSCLE TROPOMYOSINS WITH DNASE I.

L6 ANSWER 16 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI FLUORESCENCE SPECTROSCOPY OF BENZO-ALPHA-PYRENE DIOL EPOXIDE DNA ADDUCTS CONFORMATION-SPECIFIC EMISSION SPECTRA.

L6 ANSWER 17 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI CHARACTERIZATION OF THE FLUORESCENCE OF THE ANTITUMOR AGENT MITOXANTRONE.

L6 ANSWER 18 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI SPECTROFLUOROMETRIC STUDIES OF THE LIPID PROBE NILE RED.

L6 ANSWER 19 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI FLUORESCENCE OF HISTONES H-1 A TYROSINATE-LIKE FLUORESCENCE EMISSION IN CERATITIS-CAPITATA H-1 AT NEUTRAL PH VALUES.

L6 ANSWER 20 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 TI P AMINO BENZAMIDINE AS A FLUORESCENT PROBE FOR THE ACTIVE SITE OF SERINE PROTEASES.

=> d 16 ti 21-56

- L6 ANSWER 21 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on
STN
TI THE INTERACTION OF HUMAN SERUM ALBUMIN AND HEMOPEXIN WITH PORPHYRINS.
- L6 ANSWER 22 OF 56 MEDLINE on STN
TI Far-red **fluorescent** proteins evolved from a blue
chromoprotein from *Actinia equina*.
- L6 ANSWER 23 OF 56 MEDLINE on STN
TI Photophysics and biological applications of the environment-sensitive
fluorophore 6-N,N-dimethylamino-2,3-naphthalimide.
- L6 ANSWER 24 OF 56 MEDLINE on STN
TI Fluorescence and circular dichroism spectroscopic studies on bovine
lactoperoxidase.
- L6 ANSWER 25 OF 56 MEDLINE on STN
TI Salt-induced folding of alkaline denatured creatine kinase under high pH
conditions.
- L6 ANSWER 26 OF 56 MEDLINE on STN
TI Interaction of tryptophan residues of cytochrome P450_{scc} with a highly
specific fluorescence quencher, a substrate analogue, compared to
acrylamide and iodide.
- L6 ANSWER 27 OF 56 MEDLINE on STN
TI [Bertalanffy-like fluorescence staining with 3-dimethylamino-6-
methoxyacridine].
Über eine Bertalanffy-analoge Fluorochromierung mit 3-Dimethylamino-6-
methoxyacridin.
- L6 ANSWER 28 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Saturated red light-emitting copolymers of poly(aryleneethynylene)s with
narrow-band-gap (NBG) units: Synthesis and luminescent properties
- L6 ANSWER 29 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Synthesis and spectral properties of novel water-soluble near-infrared
fluorescent indocyanines
- L6 ANSWER 30 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI 7-(2-Methoxycarbonylvinyl)-3-hydroxychromones: new dyes with **red**
shifted dual emission
- L6 ANSWER 31 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI The synthesis and photophysical properties of a novel red-emitting
dioxolane-substituted pentacene derivative
- L6 ANSWER 32 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI High-Efficiency Saturated Red Emitting Polymers Derived from Fluorene and
Naphthoselenadiazole
- L6 ANSWER 33 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Facile synthesis of chlorophyll analog possessing a disulfide bond and
formation of self-assembled monolayer on gold surface
- L6 ANSWER 34 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Cd(II) Sensing in Water Using Novel Aromatic Iminodiacetate Based
Fluorescent Chemosensors
- L6 ANSWER 35 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Synthesis and Optical and Electroluminescent Properties of Novel

Conjugated Copolymers Derived from Fluorene and Benzoselenadiazole

- L6 ANSWER 36 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Development of novel cyanine dyes for diode laser induced fluorescence detection
- L6 ANSWER 37 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Synthesis and properties of two polymerizable **fluorescent** monomers
- L6 ANSWER 38 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI RGB **emission** using a dimesitylboryl-bithiophene derivative as a universal host and pentacene derivatives as the red emitters
- L6 ANSWER 39 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Far **red-shifted fluorescent** proteins and their variants from *Heteractis crispis*
- L6 ANSWER 40 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI The first genuine observation of **fluorescent** mononuclear phthalocyanine aggregates
- L6 ANSWER 41 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Probing the donor and acceptor dye assemblies at the galleries of α -zirconium phosphate
- L6 ANSWER 42 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Natural animal coloration can be determined by a nonfluorescent green **fluorescent** protein homolog
- L6 ANSWER 43 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Synthesis and properties of soluble, **fluorescent** polyesters and polyethers with substituted m-terphenyl segments in the main chain
- L6 ANSWER 44 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI The Formation and Behavior of **Fluorescent** Lewis Acid-Base Exciplexes and Triplexes
- L6 ANSWER 45 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Fluorescence Properties of Benz[f]indole, a Wavelength and Quenching Selective Tryptophan Analog
- L6 ANSWER 46 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Exploring the hydrophobic binding sites of calmodulin with **fluorescent** probes: Dynamic formation of the excimer
- L6 ANSWER 47 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Influence of Chlorination on Chromophores and Fluorophores in Humic Substances
- L6 ANSWER 48 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI An assessment of the usefulness of 5-hydroxytryptophan as an optical probe
- L6 ANSWER 49 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Protonation of free 2,9-bis(p-biphenyl)-1,10-phenanthroline sites in a 56-membered macrocycle and in its ReI and CuI complexes Absorption spectra, luminescence properties, and excited state interactions
- L6 ANSWER 50 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Aggregation state of dehydroergosterol in water and in a model system of membranes
- L6 ANSWER 51 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Fluorescence and Photodimerization Studies of Cyano-Substituted

Diphenylbutadienes

L6 ANSWER 52 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Solvent effects on the **fluorescent emission** of some
new benzimidazole derivatives

L6 ANSWER 53 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Toward red-emitting radiation tolerant chromophores

L6 ANSWER 54 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Chlorotetracycline-induced fluorescence from atherosclerotic plaque with
UV laser excitation

L6 ANSWER 55 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI α -Tocopherol as a **fluorescent** probe. 1. Its
spectroscopic characterization and use as a probe of liposome

L6 ANSWER 56 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN
TI Formation of **fluorescent** derivatives of alcohol dehydrogenase
and other proteins on UV irradiation

=> d ab bib 1,5,6,8,22,39

L13 HAS NO ANSWERS

'1 5 6 8 22 39 ' IS NOT A VALID SEARCH STATUS KEYWORD

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L13 0 SEA (FLUORESCENT OR CHROMOPROTEIN) AND EMISSION AND MAXIMUM
AND " RED SHIFTED" AND ANTHOZOAN

=> d l6 ab bib 1,5,6,8,22,39

L6 ANSWER 1 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
AB Two-photon (TP) excitation (820-1150 nm) and **emission**
(280-700 nm) spectra for the **fluorescent** proteins
(FPs) ECFP3, EGFP3 and EYFP3 produced in human tumour cells were recorded.
TP excitation spectra of pure and highly enriched samples were found to be
more differentiated in comparison with their one-photon (OP) spectra.
They exhibited more pronounced main and local **maxima**, which
coincided among different purity grades within small limits. TP and OP
emission spectra of pure and enriched samples were identical.
However, in crude samples, excitation was slightly blue-shifted and
emission red-shifted. The data indicate that
both OP and TP excitation routes led to the same excited states of these
molecules. The **emission** intensity is dependent on the pH of the
environment for both types of excitation; the **emission** intensity
maximum can be recorded in the alkaline range. Reconstitution of
emission intensity after pH quenching was incomplete, albeit that
the respective spectral profiles were identical to those prequenching.
When **emission** data were averaged over the whole range of
excitation, the resulting **emission** profile and **maximum**
coincided with the data generated by optimal excitation. Therefore,
out-of-**maximum** excitation, common practice in TP excitation
microscopy, can be used for routine application.

AN 2005:168065 BIOSIS
DN PREV200500175603
TI Two-photon excitation and **emission** spectra of the green
fluorescent protein variants ECFP, EGFP and EYFP.
AU Spiess, E. [Reprint Author]; Bestvater, F.; Heckel-Pompey, A.; Toth, K.;
Hacker, M.; Stobrawa, G.; Feurer, T.; Wotzlaw, C.; Berchner-Pfannschmidt,
U.; Porwol, T.; Acker, H.

CS Deutsch Krebsforschungszentrum, D-6900, Heidelberg, Germany
e.spiess@dkfz-heidelberg.de
SO Journal of Microscopy (Oxford), (March 2005) Vol. 217, No. 3, pp. 200-204.
print.
CODEN: JMICAR. ISSN: 0022-2720.
DT Article
LA English
ED Entered STN: 4 May 2005
Last Updated on STN: 4 May 2005

L6 ANSWER 5 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
AB We performed the biochemical and biophysical characterization of a red
fluorescent protein, eqFP611, from the sea anemone *Entacmaea*
quadricolor cloned in *Escherichia coli*. With an excitation
maximum at 559 nm and an **emission**
maximum at 611 nm, the recombinant protein shows the
most **red-shifted emission** and the largest
Stokes shift of all nonmodified proteins in the green **fluorescent**
protein family. The protein fluoresces with a high quantum yield of 0.45,
although it resembles the nonfluorescent members of this protein class, as
inferred from the absence of the key amino acid serine at position 143.
Fluorescence is constant within the range pH 4-10. Red fluorophore
maturation reaches a level of 90% after approx 12 h by passing through a
green intermediate. After complete maturation, only a small fraction of
the green species (less than 1%) persists. The protein has a reduced
tendency to oligomerize, as shown by its monomeric appearance in SDS/PAGE
analysis and single-molecule experiments. However, it forms tetramers at
higher concentrations in the absence of detergent. Fluorescence
correlation spectroscopy reveals light-driven transitions between bright
and dark states on submillisecond and millisecond time scales.
Applicability of eqFP611 for in vivo labeling in eukaryotic systems was
shown by expression in a mammalian cell culture.

AN 2002:558092 BIOSIS
DN PREV200200558092
TI A far-red **fluorescent** protein with fast maturation and reduced
oligomerization tendency from *Entacmaea quadricolor* (Anthozoa, Actinaria).
AU Wiedenmann, Joerg; Schenk, Andreas; Roecker, Carlheinz; Girod, Andreas;
Spindler, Klaus-Dieter; Nienhaus, G. Ulrich [Reprint author]
CS Department of Physics, University of Illinois at Urbana-Champaign, Urbana,
IL, 61801, USA
uli@uiuc.edu
SO Proceedings of the National Academy of Sciences of the United States of
America, (September 3, 2002) Vol. 99, No. 18, pp. 11646-11651. print.
CODEN: PNASA6. ISSN: 0027-8424.
DT Article
LA English
OS Genbank-AY130757; EMBL-AY130757; DDBJ-AY130757
ED Entered STN: 30 Oct 2002
Last Updated on STN: 30 Dec 2002

L6 ANSWER 6 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
AB The red **fluorescent** protein DsRed recently cloned from *Discosoma*
coral, with its significantly **red-shifted** excitation
and **emission maxima** (558 and 583 nm,
respectively), has attracted great interest because of its spectral
complementation to other **fluorescent** proteins, including the
green **fluorescent** protein and its enhanced mutant EGFP. We
demonstrated that the much slower DsRed fluorescence development could be
described by a three-step kinetic model, in contrast to the fast EGFP
maturation, which was fitted by a one-step model. At pH below 5.0 DsRed
fluorescence gradually decreased, and the rate and degree of this
fluorescence inactivation depended on the pH value. The kinetics of
fluorescence inactivation under acidic conditions was fitted by a
two-exponential function where the initial inactivation rate was

proportional to the fourth power of proton concentration. Subsequent DsRed alkalization resulted in partial fluorescence recovery, and the rate and degree of such recovery depended on the incubation time in the acid. Recovery kinetics had a lag-time and was fitted minimally by three exponential functions. The DsRed absorbance and circular dichroism spectra revealed that the fluorescence loss was accompanied by protein denaturation. We developed a kinetic mechanism for DsRed denaturation that includes consecutive conversion of the initial state of the protein, protonated by four hydrogen ions, to the denatured one through three intermediates. The first intermediate still emits fluorescence, and the last one is subjected to irreversible inactivation. Because of tight DsRed tetramerization we have suggested that obligatory protonation of each monomer results in the fluorescence inactivation of the whole tetramer.

AN 2002:149199 BIOSIS
 DN PREV200200149199
 TI Kinetic analysis of maturation and denaturation of DsRed, a coral-derived red fluorescent protein.
 AU Verkhusha, V. V. [Reprint author]; Akovbian, N. A.; Efremenko, E. N.; Varfolomeyev, S. D.; Vrzheschch, P. V.
 CS Center for Molecular Medicine, Lomonosov Moscow State University, Moscow, 119899, Russia
 vrzh@genebee.msu.ru
 SO Biochemistry (Moscow), (December, 2001) Vol. 66, No. 12, pp. 1342-1351. print.
 CODEN: BIORAK. ISSN: 0006-2979.
 DT Article
 LA English
 ED Entered STN: 14 Feb 2002
 Last Updated on STN: 26 Feb 2002

L6 ANSWER 8 OF 56 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
 AB DsRed, a brilliantly red fluorescent protein, was recently cloned from Discosoma coral by homology to the green fluorescent protein (GFP) from the jellyfish Aequorea. A core question in the biochemistry of DsRed is the mechanism by which the GFP-like 475-nm excitation and 500-nm emission maxima of immature DsRed are red-shifted to the 558-nm excitation and 583-nm emission maxima of mature DsRed. After digestion of mature DsRed with lysyl endopeptidase, high-resolution mass spectra of the purified chromophore-bearing peptide reveal that some of the molecules have lost 2 Da relative to the peptide analogously prepared from a mutant, K83R, that stays green. Tandem mass spectrometry indicates that the bond between the alpha-carbon and nitrogen of Gln-66 has been dehydrogenated in DsRed, extending the GFP chromophore by forming -CdbdN-CdbdO at the 2-position of the imidazolidinone. This acylimine substituent quantitatively accounts for the red shift according to quantum mechanical calculations. Reversible hydration of the CdbdN bond in the acylimine would explain why denaturation shifts mature DsRed back to a GFP-like absorbance. The CdbdN bond hydrolyses upon boiling, explaining why DsRed shows two fragment bands on SDS/PAGE. This assay suggests that conversion from green to red chromophores remains incomplete even after prolonged aging.

AN 2001:12180 BIOSIS
 DN PREV200100012180
 TI The structure of the chromophore within DsRed, a red fluorescent protein from coral.
 AU Gross, Larry A.; Baird, Geoffrey S.; Hoffman, Ross C.; Baldrige, Kim K.; Tsien, Roger Y. [Reprint author]
 CS University of California, San Diego, 9500 Gilman Drive, 310 Cellular and Molecular Medicine West 0647, La Jolla, CA, 92093-0647, USA
 SO Proceedings of the National Academy of Sciences of the United States of America, (October 24, 2000) Vol. 97, No. 22, pp. 11990-11995. print.
 CODEN: PNASA6. ISSN: 0027-8424.

DT Article
LA English
ED Entered STN: 27 Dec 2000
Last Updated on STN: 27 Dec 2000

L6 ANSWER 22 OF 56 MEDLINE on STN

AB Proteins of the GFP (green fluorescent protein) family demonstrate a great spectral and phylogenetic diversity. However, there is still an intense demand for **red-shifted** GFP-like proteins in both basic and applied science. To obtain GFP-like **chromoproteins** with **red-shifted** absorption, we performed a broad search in blue-coloured Anthozoa species. We revealed specimens of *Actinia equina* (beadlet anemone) exhibiting a bright blue circle band at the edge of the basal disc. A novel blue **chromoprotein**, aeCP597, with an absorption **maximum** at 597 nm determining the coloration of the anemone basal disk was cloned. AeCP597 carries a chromophore chemically identical with that of the well-studied DsRed (red **fluorescent** protein from *Discosoma* sp.). Thus a strong 42-nm bathochromic shift of aeCP597 absorption compared with DsRed is determined by peculiarities of chromophore environment. Site-directed and random mutagenesis of aeCP597 resulted in far-red **fluorescent** mutants with **emission maxima** at up to 663 nm. The most bright and stable mutant AQ143 possessed excitation and **emission maxima** at 595 and 655 nm respectively. Thus aeCP597 and its **fluorescent** mutants set a new record of **red-shifted** absorption and **emission maxima** among GFP-like proteins.

AN 2005640113 IN-PROCESS

DN PubMed ID: 16164420

TI Far-red **fluorescent** proteins evolved from a blue **chromoprotein** from *Actinia equina*.

AU Shkrob Maria A; Yanushevich Yuriy G; Chudakov Dmitriy M; Gurskaya Nadya G; Labas Yulii A; Poponov Sergey Y; Mudrik Nikolay N; Lukyanov Sergey; Lukyanov Konstantin A

CS Shemiakin-Ovchinnikov Institute of Bioorganic Chemistry, Miklukho-Maklaya 16/10, 117997 Moscow, Russia.

SO Biochemical journal, (2005 Dec 15) 392 (Pt 3) 649-54.
Journal code: 2984726R. ISSN: 1470-8728.

CY England: United Kingdom

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS NONMEDLINE; IN-DATA-REVIEW; IN-PROCESS; NONINDEXED; Priority Journals

ED Entered STN: 20051203

Last Updated on STN: 20051203

L6 ANSWER 39 OF 56 CAPLUS COPYRIGHT 2005 ACS on STN

AB Nucleic acid compns. encoding Stichodactylidaen **chromoproteins** and **fluorescent** mutants thereof, as well as the polypeptide compns. encoded by the same, are provided. The proteins of interest are proteins that are colored and/or **fluorescent**, where this feature arises from the interaction of two or more residues of the protein. Also of interest are proteins that are substantially similar to, or mutants of, the above specific proteins, including non-aggregating mutants and mutants with modulated oligomerization characteristics as compared to wild type. Thus, the two wild-type **chromoprotein** isoforms from *Heteractis crispa* exhibit a strong **emission max.** at .apprx.580-640 nm. Site-specific mutagenesis of the Cys-148 residue to serine dramatically increases the quantum yield of red fluorescence as compared to the wild-type protein, and further random mutagenesis (A2S, T36A, C143S, L173H, P201L, K204E) yielded an even brighter mutant. A single mutation, L126H, may be responsible for modifying the oligomeric state of the protein from tetrameric to dimeric. Also provided are fragments of the nucleic acids and the peptides encoded

thereby, as well as antibodies to the subject proteins and transgenic cells and organisms. The subject protein and nucleic acid compns. find use in a variety of different applications. Finally, kits for use in such applications, e.g., that include the subject nucleic acid compns., are provided.

AN 2002:293685 CAPLUS

DN 136:321045

TI Far **red-shifted fluorescent** proteins and their variants from *Heteractis crispis*

IN Lukyanov, Sergey Anatolievich; Fradkov, Arcady Fedorovich; Lukyanov, Konstantin Anatolievich; Gurskaya, Nadezda Georgievna

PA Clontech Laboratories, Inc., USA

SO PCT Int. Appl., 81 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 17

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002030965	A2	20020418	WO 2001-US32080	20011012
	WO 2002030965	A3	20030605		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	CA 2425318	AA	20020418	CA 2001-2425318	20011012
	AU 2002011722	A5	20020422	AU 2002-11722	20011012
	EP 1334122	A2	20030813	EP 2001-979797	20011012
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
	JP 2004532606	T2	20041028	JP 2002-534350	20011012
PRAI	US 2000-240018P	P	20001012		
	US 2001-306131P	P	20010716		
	WO 2001-US32080	W	20011012		